**SEQUENCE LISTING** 

**GENERAL INFORMATION:** 

(i)

APPLICANT: PEREGRINO FERREIRA, Paulo;

5 GESSIEN KROON, Erna;

PIMENTA DOS REIS, Karlisson Jennner;

BIAS FORTES FERRAZ, Isabella;

CERQUEIRA LEITE, Romulo.

(ii)

TITLE OF INVENTION: Method and composition for the diagnosis of equine infectious anemia virus disease by using the recombinant capsid protein virus (p26)

(iii)

**NUMBER OF SEQUENCES: 1** 

15 (iv)

**CORRESPONDENCE ADDRESS:** 

(A)

ADDRESSEE: Universidade Federal de Minas Gerais - CTIT

(B)

20 STREET: Avenida Antônio Carlos, 6627 Bairro São Francisco

(C)

CITY: Belo Horizonte

(D)

STATE: Minas Gerais

25 **(E)** 

COUNTRY: BRAZIL

(F)

ZIP: 31270-901

(v)

30 COMPUTER READABLE FORM:

(A)

MEDIUM TYPE: diskette - 3.50 inch, 1.44 Mb storage

(B)

COMPUTER: IBM compatible

(C)

5 OPERATING SYSTEM: Windows 98

(D)

SOFTWARE: Office premium

(vi)

**CURRENT APPLICATION DATA:** 

10 (A)

APPLICATION NUMBER: U.S. 09/331.262

(B)

**FILING DATE:** 

(C)

15 CLASSIFICATION: C12Q1/70

(vii)

PRIOR APPLICATION DATA

(A)

APPLICATION NUMBER: PI 9606273-8

20 (B)

FILING DATE: 18-DEC-1996

(2)

INFORMATION FOR SEQ ID N0:1:

(i)

25 SEQUENCE CHARACTERISTICS:

(A)

LENGHT: 252 amino acids

(B)

TYPE: amino acid

30 (D)

TOPOLOGY: linear

```
(ii)
    MOLECULE TYPE: protein
    (vi)
    ORIGINAL SOURCE
    (A)
    ORGANISM : equine infectious anemia virus
    (ix)
    FEATURE:
    (A)
10
    NAME: p26
    (x)
    PUBLICATION INFORMATION
    (A)
    AUTHORS:
15
   . (B)
    TITLE: (
    C)
    JOURNAL:
    (D)
20
    VOLUME:
    (F)
    PAGES:
    (G)
    DATE:
25
    (xi)
    SEQUENCE DESCRIPTION: SEQ ID NO:1
    His His His His His Gly Ser Pro Gly Asn Pro Leu Thr Trp
                  5
                                   10
                                                     15
30
```

	Ser Lys Ala Leu Lys Lys Leu Glu Lys Val Thr Val Gln Gly			
	20	25	30	
	Gln Lys Leu Thr Thr Gly A	Asn Cys Na Trp Ala Leu Ser	Leu Val	
	35	40	45	
5	Asp Leu Phe His Asp Thr Asn Phe Val Lys Glu Lys Asp Trp Gln			
	50	55	60	
	Leu Arg Asp Val Ile Pro Le	eu Leu Glu Asp Val Thr Gln	Thr Val	
	65	70	· . 75	
	Ser Gly Gln Glu Arg Glu A	Ala Phe Glu Arg Thr Trp Trp	Ala Ile	
10	80	85	90.	
	Ser Ala Val Lys Met Gly L	eu Gin Ile Asn AsnVal Val <i>A</i>	Asp Gly	
	95	100	105	
	Lys Ala Ser Phe Gln Leu I	Leu Arg Ala Lys Tyr Glu Lys	Lys Thr	
	110	115	120	
15	Ala Asn Lys Lys Gln Ser (	Glu Pro Ser Glu Glu Tyr Pro	lle Met	
	125	130	135	
	lle Asp Gly Ala Gly Asn A	rg Asn Phe Arg Pro Leu Thi	Pro Arg	
	140	145	150	
	Gly Tyr Thr Thr Trp Val As	snThr lle Gln Thr Asn Gly L	eu Leu	
20	155	160	165	
	Asn Glu Ala Ser Gln Asn I	Leu Phe Gly Ile Leu Ser Val	Asp Cys	
	170	175	180	
	Thr Ser Glu Glu Met Asn	Ala Phe Leu Asp Val Val Pr	o Gly Gln	
	185	190	195	
25		eu Leu Asp Ala Ile Asp Lys I		
	200	205	210	
		His Pro Leu Pro Asn Ala P	ro Leu Val	
•	215	220	225	
		e Pro Met Thr Ala Arg Phe	_	
30	230	235	240	
	Gly Leu Gly Val Pro Arg G	_	•	
	245	250		

Asn Cys Val Val Gln Ser Phe Gly Val Ile Gly Gln Ala His Leu.			
260	265	270	
Glu Leu Pro Arg Pro Asn Lys Arg Ile Arg Asn Gln. Ser Phe As			
275	280	285	
Gln Tyr Asn Cys Ser lle Asn. Asn Lys Thr Glu Leu Glu Thr Trp			
290	295	300	
Lys Leu.Val Lys Thr Ser Gly Val Thr Pro Leu Pro. lle Ser Ser			
305	310	315	
Glu Ala Asn Thr Gly Leu	*		
320			